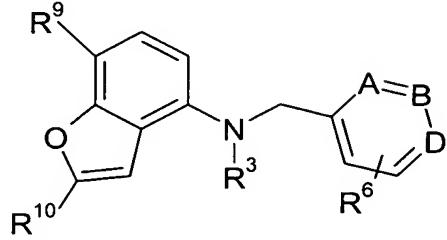
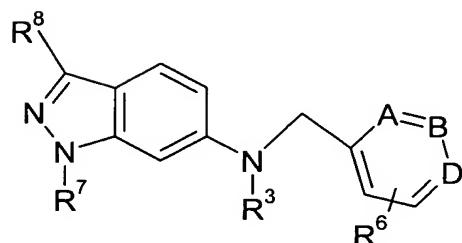
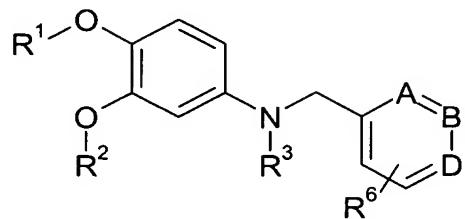


This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Previously Presented): An isolated compound of Formulas I-III:



wherein

one of A, B and D is N-O and the others are CR⁶;

R^1 is alkyl having 1 to 4 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen;

R^2 is alkyl having 1 to 12 carbon atoms which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, hydroxy, cyano, C_1 -4-alkoxy, oxo or combinations thereof, and wherein optionally one or more - CH_2CH_2 - groups is replaced in each case by $-CH=CH-$ or $-C\equiv C-$,

cycloalkyl having 3 to 10 carbon atoms which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, or combinations thereof,

cycloalkylalkyl having 4 to 16 carbon atoms which is unsubstituted or substituted in the cycloalkyl portion and/or the alkyl portion one or more times by halogen, oxo, cyano, hydroxy, C_1 -4-alkyl, C_1 -4-alkoxy or combinations thereof,

aryl having 6 to 14 carbon atoms, which is unsubstituted or substituted one or more times by halogen, CF_3 , OCF_3 , alkyl, hydroxy, alkoxy, nitro, methylenedioxy, ethylenedioxy, cyano, or combinations thereof,

arylalkyl in which the aryl portion has 6 to 14 carbon atoms and the alkyl portion, which is branched or unbranched, has 1 to 5 carbon atoms, wherein the arylalkyl radical is unsubstituted or is substituted in the aryl portion one or more times by halogen, CF_3 , OCF_3 , alkyl, hydroxy, alkoxy, nitro, cyano, methylenedioxy, ethylenedioxy, or combinations thereof, and wherein in the alkyl portion one or more - CH_2CH_2 - groups are each optionally replaced by $-CH=CH-$ or $-C\equiv C-$, and/or one or more - CH_2 - groups are each optionally replaced by $-O-$ or $-NH-$ and/or the alkyl portion is optionally substituted by halogen, oxo, hydroxy, cyano, or combinations thereof,

a partially unsaturated carbocyclic group having 5 to 14 carbon atoms, which is unsubstituted or substituted one or more times by halogen, alkyl, alkoxy,

hydroxy, nitro, cyano, oxo, or combinations thereof,

a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is an N, O or S atom, wherein the heterocyclic group is unsubstituted or substituted one or more times by halogen, hydroxy, aryl, alkyl, alkoxy, cyano, trifluoromethyl, nitro, oxo, or combinations thereof, or

a heterocycle-alkyl group, wherein the heterocyclic portion is saturated, partially saturated or unsaturated, and has 5 to 10 ring atoms in which at least 1 ring atom is an N, O or S atom, and the alkyl portion is branched or unbranched and has 1 to 5 carbon atoms, the heterocycle-alkyl group is unsubstituted or substituted one or more times in the heterocyclic portion by halogen, OCF₃, hydroxy, aryl, alkyl, alkoxy, cyano, trifluoromethyl, nitro, oxo, or combinations thereof, wherein in the alkyl portion one or more -CH₂CH₂- groups are each optionally replaced by -CH=CH- or -C≡C-, and/or one or more -CH₂- groups are each optionally replaced by -O- or -NH- and/or the alkyl portion is optionally substituted by halogen, oxo, hydroxy, cyano, or combinations thereof;

R³ is H,

cycloalkyl having 3 to 10 carbon atoms which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, or combinations thereof,

aryl having 6 to 14 carbon atoms and which is unsubstituted or substituted one or more times by halogen, alkyl, alkenyl, alkynyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, OCF₃, amino, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, pyrrolyl, tetrazole-5-

yl, 2(-heterocycle)tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, trialkylsilyloxy, R^4 -L-, or combinations thereof,

heteroaryl having 5 to 10 ring atoms in which at least 1 ring atom is a heteroatom, which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, trialkylsilyloxy, R^4 -L-, or combinations thereof,

a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is an N, O or S atom, which is unsubstituted or substituted one or more times by halogen, hydroxy, aryl, alkyl, alkoxy, cyano, trifluoromethyl, nitro, oxo, or combinations thereof;

R^4 is H,

alkyl having 1 to 8 carbon atoms which is unsubstituted or substituted one or more times by halogen, C_{1-4} -alkyl, C_{1-4} -alkoxy, oxo, or combinations thereof,

alkylamino or dialkylamino wherein each alkyl portion has independently 1 to 8 carbon atoms,

a partially unsaturated carbocycle-alkyl group wherein the carbocyclic portion has 5 to 14 carbon atoms and the alkyl portion has 1 to 5 carbon atoms, which is unsubstituted or substituted one or more times by halogen, alkyl, alkoxy, nitro, cyano, oxo, or combinations thereof,

cycloalkyl having 3 to 10 carbon atoms which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkoxy, alkyl having 1 to 4 carbon atoms, or combinations thereof,

cycloalkylalkyl having 4 to 16 carbon atoms which is unsubstituted or substituted in the cycloalkyl portion and/or the alkyl portion one or more times by halogen, oxo, cyano, hydroxy, alkyl, alkoxy or combinations thereof,

aryl having 6 to 14 carbon atoms and which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy, dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxycarbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, cycloalkyl, aryl, heteroaryl, or combinations thereof,

arylalkyl having 7 to 19 carbon atoms, wherein the aryl portion has 6 to 14 carbon atoms and the alkyl portion, which is branched or unbranched, has 1 to 5 carbon atoms, wherein the arylalkyl radical is unsubstituted or substituted, in the aryl portion, one or more times by halogen, trifluoromethyl, CF_3O , nitro, amino, alkyl, alkoxy, amino, alkylamino, dialkylamino, or combinations thereof, and/or substituted in the alkyl portion by halogen, cyano, methyl, or combinations thereof, wherein in the alkyl portion one or more $-\text{CH}_2\text{CH}_2-$ groups are each optionally replaced by $-\text{CH}=\text{CH}-$ or $-\text{C}\equiv\text{C}-$, and/or one or more $-\text{CH}_2-$ groups are each optionally replaced by $-\text{O}-$ or $-\text{NH}-$,

a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is an N, O or S atom, which is

unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, or combinations thereof, or

a heterocycle-alkyl group, wherein the heterocyclic portion is saturated, partially saturated or unsaturated, and has 5 to 10 ring atoms in which at least 1 ring atom is a N, O or S atom, and the alkyl portion which is branched or unbranched and has 1 to 5 carbon atoms, the heterocycle-alkyl group is unsubstituted or substituted one or more times in the heterocyclic portion by halogen, alkyl, alkoxy, cyano, trifluoromethyl, CF_3O , nitro, oxo, amino, alkylamino, dialkylamino, or combinations thereof and/or substituted in the alkyl portion by halogen, cyano, or methyl or combinations thereof;

L is a single bond or a divalent aliphatic radical having 1 to 8 carbon atoms wherein one or more $-\text{CH}_2-$ groups are each optionally replaced by $-\text{O}-$, $-\text{S}-$, $-\text{SO}-$, $-\text{SO}_2-$, $-\text{NR}^5-$, $-\text{SO}_2\text{NH}-$, $-\text{NHSO}_2-$, $-\text{SO}_2\text{NR}^5-$, $-\text{NR}^5\text{SO}_2-$, $-\text{CO}-$, $-\text{NR}^5\text{CO}-$, $-\text{CONR}^5-$, $-\text{NHCONH}-$, $-\text{OCONH}-$, $-\text{NHCOO}-$, $-\text{SCONH}-$, $-\text{SCSNH}-$, or $-\text{NHCSNH}-$;

R^5 is H,

alkyl having 1 to 8 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, C_{1-4} -alkyl, C_{1-4} -alkoxy, oxo, or combinations thereof,

aryl having 6 to 14 carbon atoms and which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, ethylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, or combinations thereof, or

arylalkyl having 7 to 19 carbon atoms, wherein the aryl portion has 6 to 14 carbon atoms and the alkyl portion, which is branched or unbranched, has 1 to 5 carbon atoms, wherein the arylalkyl radical is unsubstituted or substituted, in the aryl portion, one or more times by halogen, trifluoromethyl, CF_3O , nitro, amino, alkyl, alkoxy, amino, alkylamino, dialkylamino, or combinations thereof, and/or substituted in the alkyl portion by halogen, cyano, methyl, or combinations thereof, wherein in the alkyl portion one or more $-\text{CH}_2\text{CH}_2-$ groups are each optionally replaced by $-\text{CH}=\text{CH}-$ or $-\text{C}\equiv\text{C}-$, and/or one or more $-\text{CH}_2-$ groups are each optionally replaced by $-\text{O}-$ or $-\text{NH}-$;

R^6 is H, halogen, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, CN, or hydroxyl;

R^7 is H,

alkyl having 1 to 8 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, cyano, hydroxy, C_{1-4} - alkoxy, or combinations thereof

cycloalkyl having 3 to 10 carbon atoms which is unsubstituted or substituted one

or more times by halogen, hydroxy, oxo, cyano, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, or combinations thereof,

cycloalkylalkyl having 4 to 16 carbon atoms which is unsubstituted or substituted in the cycloalkyl portion and/or the alkyl portion one or more times by halogen, oxo, cyano, hydroxy, C₁₋₄-alkyl, C₁₋₄-alkoxy or combinations thereof,

aryl having 6 to 14 carbon atoms, which is unsubstituted or substituted one or more times by halogen, CF₃, OCF₃, alkyl, hydroxy, alkoxy, nitro, methylenedioxy, ethylenedioxy, cyano, or combinations thereof,

arylalkyl in which the aryl portion has 6 to 14 carbon atoms and the alkyl portion, which is branched or unbranched, has 1 to 5 carbon atoms, wherein the arylalkyl radical is unsubstituted or is substituted in the aryl portion one or more times by halogen, CF₃, OCF₃, alkyl, hydroxy, alkoxy, nitro, cyano, methylenedioxy, ethylenedioxy, or combinations thereof, and wherein in the alkyl portion one or more -CH₂CH₂- groups are each optionally replaced by -CH=CH- or -C≡C-, and/or one or more -CH₂- groups are each optionally replaced by -O- or -NH- and/or the alkyl portion is optionally substituted by halogen, oxo, hydroxy, cyano, or combinations thereof,

a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is an N, O or S atom, which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, or combinations thereof, or

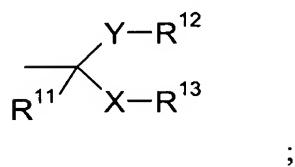
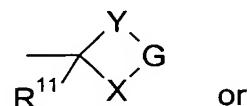
a heterocycle-alkyl group, wherein the heterocyclic portion is saturated, partially saturated or unsaturated, and has 5 to 10 ring atoms in which at least 1 ring atom is a N, O or S atom, and the alkyl portion which is branched or unbranched and has 1 to 5 carbon atoms, the heterocycle-alkyl group is unsubstituted or substituted one or more times in the heterocyclic portion by halogen, alkyl, alkoxy, cyano, trifluoromethyl, CF_3O , nitro, oxo, amino, alkylamino, dialkylamino, or combinations thereof and/or substituted in the alkyl portion by halogen, cyano, or methyl or combinations thereof;

R^8 is H, or

alkyl having 1 to 4 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, cyano, and/or C_{1-4} - alkoxy, and one or more $-\text{CH}_2\text{CH}_2-$ groups can be replaced in each case by $-\text{CH}=\text{CH}-$ or $-\text{C}\equiv\text{C}-$;

R^9 is alkoxy or alkylthio, in each case having 1 to 4 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen;

R^{10} is $-\text{CO-C}_{1-4}\text{-alkyl}$ which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, or is



R¹¹ is H or alkyl having 1 to 4 carbon atoms, which is branched or unbranched, and which is unsubstituted or substituted one or more times by halogen;

R¹² is alkyl having 1 to 6 carbon atoms, which is branched or unbranched, and which is unsubstituted or substituted one or more times by halogen;

R¹³ is alkyl having 1 to 6 carbon atoms, which is branched or unbranched, and which is unsubstituted or substituted one or more times by halogen;

X and Y are each independently O or S; and

G is alkylene having 2 to 7 carbon atoms which is unsubstituted or substituted one or more times by halogen; or

a pharmaceutically acceptable salt thereof;

wherein an optically active compound can be in the form of one of its separate enantiomers or mixtures thereof, including racemic mixtures.

2. (Previously Presented): An isolated compound according to claim 1, wherein B is N-O.

3. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula I and R¹ is methyl or difluoromethyl.

4. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula I and R² is cycloalkyl.

5. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula I and R² is phenyl, methylphenyl, methoxyphenyl, chlorophenyl, phenethyl, phenpropyl, phenbutyl, phenylethenyl, phenoxyethyl, phenoxypropyl, phenoxybutyl, chlorophenylethyl, methoxyphenylethyl, chlorophenylethenyl, chlorophenoxyethyl, chlorophenylpropyl, methoxyphenpropyl, methoxyphenbutyl, chlorophenbutyl, nitrophenbutyl, or chlorophenylaminoethyl.

6. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula I and R² is cyclohexenyl, cyclohexadienyl, or indan-2-yl.

7. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula I and R² is methyl, difluoromethyl, trifluoromethyl, or methoxyethyl.

8. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula I and R² is tetrahydrofuranyl, pyrrolidinyl, pyrrolyl, pyridylmethyl, pyridylethyl, pyridylpropyl, piperazinylmethyl, piperazinylethyl, or methylpiperazinylethyl.

9. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula I and R² cyclopentyl, tetrahydrofuran-2-yl, CHF₂, methoxyethyl, cyclopropylmethyl, phenethyl, phenpropyl, phenylethenyl, phenoxyethyl, phenoxybutyl, phenylaminoethyl, indan-2-yl, pyridylethyl, or pyridylpropyl.

10. (Previously Presented): An isolated compound according to claim 1, wherein R³ is phenyl, naphthyl, biphenyl, furanyl, pyrazinyl, pyrimidinyl, pyridyl, quinolinyl, or isoquinolinyl, which in each case is unsubstituted or is substituted one or more times.

11. (Previously Presented): An isolated compound according to claim 10, wherein R³ is substituted by OH, F, Cl, CF₃, methyl, ethyl, methoxy, ethoxy, CN, vinyl, CH₂OH, CONHOH, CONH₂, methylenedioxy, COOH, or combinations thereof.

12. (Previously Presented): An isolated compound according to claim 1, wherein R^3 is pyridyl or phenyl which in each case is substituted or unsubstituted.

13. (Previously Presented): An isolated compound according to claim 1, wherein R^3 is phenyl substituted by halogen, COOH and/or CN.

14. (Previously Presented): An isolated compound according to claim 1, wherein R^3 is aryl substituted by R^4 -, R^4 -O-, R^4 -CO-, R^4 -NH-CO-, R^4 -SO₂-NH-, R^4 -SO₂-NHCO-, R^4 -SO₂-NH-alkylene-O-, NH₂-alkyl-NH-CO-, R^4 -alkylene-NH-CO-, alkyl-CO-NH-alkyl-, methyl, ethyl, Cl, F, CN, OCH₃, CF₃, amino, nitro, CH₂OH or COOH.

15. (Previously Presented): An isolated compound according to claim 1, wherein R^3 is phenyl substituted by R^4 -SO₂-NH- and R^4 is methyl, ethyl, propyl or phenyl.

16. (Previously Presented): An isolated compound according to claim 1, wherein R^3 is phenyl substituted by R^4 -SO₂-NH-alkylene-O-, R^4 is methyl, ethyl, propyl or phenyl, and alkylene is -CH₂-, -CH₂CH₂- or -CH₂CH₂CH₂-.

17. (Previously Presented): An isolated compound according to claim 1, wherein R^3 is phenyl substituted by R^4 -L-, R^4 is phenyl, tetrazolyl, oxazinyl, piperazinyl, methylpiperazinyl, pyridyl, methylpyridyl, pyrrolinyl, methylpyrrolinyl, piperadinyl, or methylpiperadinyl, and L is a single bond, -O-, -CO-, -CH₂-, -CH₂CH₂-, -CH₂CH₂CH₂-, -CH₂-O-, -CH₂CH₂-O-, -CH₂CH₂CH₂-O-, -CH₂-NH-CH₂CH₂-O-, -CO-NH-, -NH-CO-, or -CONHSO₂-.

18. (Previously Presented): An isolated compound according to claim 1, wherein R^6 is H or F.

19. (Previously Presented): An isolated compound according to claim 1, wherein R⁶ is H.

20. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula II and R⁷ is alkyl having 2 to 4 carbon atoms which is optionally substituted by halogen.

21. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula II and R⁷ is cyclopentyl or cyclohexyl.

22. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula II and R⁸ is H or C₂H₅.

23. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula III and R⁹ is CH₃ or C₂H₅.

24. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula III and R¹⁰ is -CO-C₁₋₄-alkyl.

25. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula III and R¹¹ is -CH₃.

26. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula III and R¹² and R¹³ are each independently -CH₃ or -CH₂CH₃.

27. (Previously Presented): An isolated compound according to claim 1, wherein X and Y are each O.

28. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula III and G is -CH₂CH₂-.

29. (Previously Presented): An isolated compound according to claim 1, wherein R³ is H.

30. (Previously Presented): An isolated compound according to claim 1, wherein D is N-O.

31. (Previously Presented): An isolated compound according to claim 1, wherein:
each aryl group is, independently, a phenyl, naphthyl or biphenyl group optionally substituted one or more times by halogen, alkyl, hydroxy, alkoxy, nitro, methylenedioxy, ethylenedioxy, amino, alkylamino, dialkylamino, hydroxyalkyl, hydroxyalkoxy, carboxy, cyano, acyl, alkoxycarbonyl, alkylthio, alkylsulfinyl, alkylsulfonyl, or phenoxy;

each heteroaryl group is, independently, a furyl, thienyl, pyrrolyl, pyrazolyl, imidazolyl, triazolyl, tetrazolyl, dithialyl, oxathialyl, isoxazolyl, oxazolyl, thiazolyl, isothiazolyl, oxadiazolyl, oxatriazolyl, dioxazolyl, oxathiazolyl, thiadiazolyl, pyridyl, pyridazinyl, pyrimidinyl, pyrazinyl, triazinyl, oxazinyl, isoxazinyl, oxathiazinyl, oxadiazinyl, benzofuranyl, isobenzofuranyl, thionaphthienyl, isothionaphthienyl, indolyl, isoindolyl, indazolyl, benzisoxazolyl, benzoxazolyl, benzthiazolyl, benzisothiazolyl, purinyl, benzopyranyl, quinolinyl, isoquinolinyl, cinnolinyl, quinazolinyl, naphthyridinyl, or benzoxazinyl group optionally substituted in one or more places by halogen, aryl, alkyl, alkoxy, carboxy, methylene, cyano, trifluoromethyl, nitro, oxo, amino, alkylamino, or dialkylamino; and

each heterocycle group is, independently, a heteroaryl group or a tetrahydrofuranyl, piperidinyl, or pyrrolidinyl group optionally substituted in one or more places by halogen, aryl, alkyl, alkoxy, carboxy, methylene, cyano, trifluoromethyl, nitro, oxo, amino, alkylamino, or dialkylamino.

32. (Previously Presented): An isolated compound according to claim 1, wherein R¹

is methyl or CHF_2 ; R^2 is alkyl, alkenyl, alkynyl, cycloalkyl, arylalkyl, heterocycle-alkyl, cycloalkylalkyl, aryl, or heterocyclic, in each case substituted or unsubstituted; and R^3 is aryl or heteroaryl, in each case substituted or unsubstituted.

33. (Previously Presented): An isolated compound according to claim 1, wherein R^1 is methyl or CHF_2 ; and R^2 is cyclopentyl, CHF_2 , cyclopropylmethyl, pyridylethyl, or tetrahydrofuranyl.

34. (Previously Presented): An isolated compound according to claim 1, wherein R^1 is methyl or CHF_2 ; R^2 is cyclopentyl, CHF_2 , cyclopropylmethyl, pyridylethyl, or tetrahydrofuranyl; and R^3 is aryl or heteroaryl, in each case substituted or unsubstituted.

35. (Previously Presented): An isolated compound according to claim 1, wherein R^1 is methyl or CHF_2 ; R^2 is cyclopentyl; and R^3 is substituted or unsubstituted aryl or heteroaryl.

36. (Previously Presented): An isolated compound according to claim 1, wherein R^1 is methyl; R^2 is cyclopentyl; and R^3 is phenyl which is substituted or unsubstituted.

37. (Previously Presented): An isolated compound according to claim 1, wherein R^1 is methyl; R^2 is cyclopentyl; and R^3 is phenyl or phenyl substituted with 1 to 3 substituents.

38. (Previously Presented): An isolated compound according to claim 1, wherein R^1 is methyl; R^2 is cyclopentyl; and R^3 is phenyl, naphthyl, biphenyl, pyridyl, pyrimidinyl, thiazolyl, pyrazinyl, quinolinyl, or isoquinolinyl, in each case substituted or unsubstituted.

39. (Previously Presented): An isolated compound according to claim 1, wherein R^1 is methyl or CHF_2 .

40. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, and B is N-O.

41. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, and R² is cyclopentyl, CHF₂, cyclopropylmethyl, pyridylethyl, or tetrahydrofuranyl.

42. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, B is N-O, and R² is cyclopentyl, CHF₂, cyclopropylmethyl, pyridylethyl, or tetrahydrofuranyl.

43. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, and R³ is 3-pyridyl or phenyl, which in each case is substituted or unsubstituted.

44. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, B is N-O, and R³ is 3-pyridyl or phenyl, which in each case is substituted or unsubstituted.

45. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, R² is cyclopentyl, CHF₂, cyclopropylmethyl, pyridylethyl or tetrahydrofuranyl, and R³ is 3-pyridyl or phenyl, which in each case is substituted or unsubstituted.

46. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, B is N-O, R² is cyclopentyl, CHF₂, cyclopropylmethyl, pyridylethyl, or tetrahydrofuranyl, and R³ is 3-pyridyl or phenyl, which in each case is substituted or unsubstituted.

47. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, and R³ is phenyl which is substituted in the 3- or 4- position.

48. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, B is N-O, and R³ is phenyl which is substituted in the 3- or 4- position.

49. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, R² is cyclopentyl, CHF₂, cyclopropylmethyl, pyridylethyl, or tetrahydrofuranyl, and R³ is phenyl which is substituted in the 3- or 4- position.

50. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, B is N-O, R² is cyclopentyl, CHF₂, cyclopropylmethyl, pyridylethyl, or tetrahydrofuranyl, and R³ is phenyl which is substituted in the 3- or 4- position.

51. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, and R³ is 3-pyridyl, 3-COOH-phenyl, 3-Cl-phenyl, 3-cyano-phenyl, 3-ethylsulfonamido-phenyl, 3-tetrazol-5-yl-phenyl, 3-hydroxymethyl-phenyl, 4-pyridyl, 4-COOH-phenyl, 4-cyano-phenyl, 4-ethylsulfonamido-phenyl, 4-tetrazol-5-yl-phenyl, or 4-hydroxymethyl-phenyl.

52. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, B is N-O, and R³ is 3-pyridyl, 3-COOH-phenyl, 3-Cl-phenyl, 3-cyano-phenyl, 3-ethylsulfonamido-phenyl, 3-tetrazol-5-yl-phenyl, 3-hydroxymethyl-phenyl, 4-pyridyl, 4-COOH-phenyl, 4-cyano-phenyl, 4-ethylsulfonamido-phenyl, 4-tetrazol-5-yl-phenyl, or 4-hydroxymethyl-phenyl.

53. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, R² is cyclopentyl, CHF₂, cyclopropylmethyl, pyridylethyl, or tetrahydrofuranyl, and R³ is 3-pyridyl, 3-COOH-phenyl, 3-Cl-phenyl, 3-cyano-phenyl, 3-ethylsulfonamido-phenyl, 3-tetrazol-5-yl-phenyl, 3-hydroxymethyl-phenyl, 4-pyridyl, 4-COOH-phenyl, 4-cyano-phenyl, 4-ethylsulfonamido-phenyl, 4-tetrazol-5-yl-phenyl, or 4-hydroxymethyl-phenyl.

54. (Previously Presented): An isolated compound according to claim 1, wherein R¹ is methyl or CHF₂, B is N-O, R² is cyclopentyl, CHF₂, cyclopropylmethyl, pyridylethyl, or tetrahydrofuranyl, and R³ is 3-pyridyl, 3-COOH-phenyl, 3-Cl-phenyl, 3-cyano-phenyl, 3-ethylsulfonamido-phenyl, 3-tetrazol-5-yl-phenyl, 3-hydroxymethyl-phenyl, 3-nitro-phenyl, 4-pyridyl, 4-COOH-phenyl, 4-cyano-phenyl, 4-ethylsulfonamido-phenyl, 4-tetrazol-5-yl-phenyl, or 4-hydroxymethyl-phenyl.

55. (Previously Presented): An isolated compound according to claim 1, wherein R³ is H or is aryl or heteroaryl, in each case substituted or unsubstituted.

56. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula II, R⁷ is cycloalkyl; and R⁸ is H or C₂H₅.

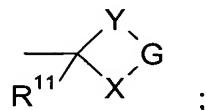
57. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula II, R⁷ is cycloalkyl; R⁸ is H or C₂H₅; and R³ is H or is aryl or heteroaryl, in each case substituted or unsubstituted.

58. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula II, R⁷ is cyclopentyl; R⁸ is H or C₂H₅; and R³ is H or is aryl or heteroaryl, in each case substituted or unsubstituted.

59. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula II, R⁷ is cyclopentyl; R⁸ is H or C₂H₅; and R³ is phenyl which is substituted or unsubstituted.

60. (Previously Presented): An isolated compound according to claim 1, wherein R³ is H or is aryl or heteroaryl, in each case substituted or unsubstituted.

61. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula III, R⁹ is alkoxy having 1 to 4 carbon atoms; R¹⁰ is COCH₃ or

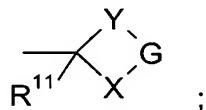


R¹¹ is -CH₃;

X and Y are both O or S; and

G is -CH₂CH₂-.

62. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula III, R³ is H or is aryl or heteroaryl, in each case substituted or unsubstituted; R⁹ is alkoxy having 1 to 4 carbon atoms; R¹⁰ is COCH₃ or



R¹¹ is -CH₃;

X and Y are both O or S; and

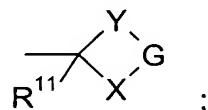
G is -CH₂CH₂-.

63. (Previously Presented): An isolated compound according to claim 1, wherein said compound is of Formula III and

R^3 is phenyl which is substituted or unsubstituted;

R^9 is alkoxy having 1 to 4 carbon atoms;

R^{10} is $COCH_3$ or



R^{11} is $-CH_3$;

X and Y are both O or S; and

G is $-CH_2CH_2-$.

64. (Previously Presented): An isolated compound according to claim 1, wherein

B is $N-O$;

R^1 is methyl or difluoromethyl;

R^2 is phenyl, methylphenyl, methoxyphenyl, chlorophenyl, phenethyl, phenpropyl, phenbutyl, phenylethenyl, phenoxyethyl, phenoxypropyl, phenoxybutyl, chlorophenylethyl, methoxyphenylethyl, chlorophenylethenyl, chlorophenoxyethyl, chlorophenylpropyl, methoxyphenylpropyl, methoxyphenbutyl, chlorophenbutyl, nitrophenbutyl, chlorophenylaminoethyl, cyclohexenyl, cyclohexadienyl, indan-2-yl methyl, difluoromethyl, trifluoromethyl, methoxyethyl, tetrahydrofuranyl, pyrrolidinyl, pyrrolyl, pyridylmethyl, pyridylethyl, pyridylpropyl, piperazinylmethyl, piperazinylethyl, methylpiperazinylethyl, cyclopentyl, CHF_2 , methoxyethyl, cyclopropylmethyl, or phenylaminoethyl;

R^3 is phenyl or pyridyl, which in each case is unsubstituted or substituted;

R^6 is H or F;

R^7 is cyclopentyl, cyclohexyl, or alkyl having 2 to 4 carbon atoms which is optionally substituted by halogen,

R^8 H or C_2H_5 ;

R^9 is CH_3 or C_2H_5 ;

R¹⁰ is -CO-C₁₋₄-alkyl;

R¹¹ is -CH₃;

R¹² and R¹³ are each independently -CH₃ or -CH₂CH₃;

X and Y are each O; and

G is -CH₂CH₂-.

65. (Previously Presented): An isolated compound according to claim 1, wherein one of A, B and D is N-O and the others are CH;

R¹ is alkyl having 1 to 4 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen;

R² is alkyl having 1 to 12 which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, hydroxy, cyano, C₁₋₄-alkoxy, oxo or combinations thereof, and wherein optionally one or more -CH₂CH₂- groups is replaced in each case by -CH=CH- or -C≡C-,

cycloalkyl having 3 to 10 carbon atoms which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, or combinations thereof,

cycloalkylalkyl having 4 to 16 carbon atoms which is unsubstituted or substituted in the cycloalkyl portion and/or the alkyl portion one or more times by halogen, oxo, cyano, hydroxy, C₁₋₄-alkyl, C₁₋₄-alkoxy or combinations thereof,

aryl having 6 to 14 carbon atoms, which is unsubstituted or substituted one or more times by halogen, CF₃, OCF₃, alkyl, hydroxy, alkoxy, nitro, methylenedioxy,

ethylenedioxy, cyano, or combinations thereof,

arylalkyl in which the aryl portion has 6 to 14 carbon atoms and the alkyl portion, which is branched or unbranched, has 1 to 5 carbon atoms, which the arylalkyl radical is unsubstituted or is substituted in the aryl portion one or more times by halogen, CF_3 , OCF_3 , alkyl, hydroxy, alkoxy, nitro, cyano, methylenedioxy, ethylenedioxy, or combinations thereof, and wherein in the alkyl portion one or more $-\text{CH}_2\text{CH}_2-$ groups are each optionally replaced by $-\text{CH}=\text{CH}-$ or $-\text{C}\equiv\text{C}-$, and one or more $-\text{CH}_2-$ groups are each optionally replaced by $-\text{O}-$ or $-\text{NH}-$ and/or the alkyl portion is optionally substituted by halogen, oxo, hydroxy, cyano, or combinations thereof,

a partially unsaturated carbocyclic group having 5 to 14 carbon atoms, which is unsubstituted or substituted one or more times by halogen, alkyl, alkoxy, hydroxy, nitro, cyano, oxo, or combinations thereof,

a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is a N, O or S atom, which is unsubstituted or substituted one or more times by halogen, hydroxy, aryl, alkyl, alkoxy, cyano, trifluoromethyl, nitro, oxo, or combinations thereof, or

a heterocycle-alkyl group, wherein the heterocyclic portion is saturated, partially saturated or unsaturated, and has 5 to 10 ring atoms in which at least 1 ring atom is a N, O or S atom, and the alkyl portion is branched or unbranched and has 1 to 5 carbon atoms, the heterocycle-alkyl group is unsubstituted or substituted one or more times in the heterocyclic portion by halogen, OCF_3 , hydroxy, aryl, alkyl, alkoxy, cyano, trifluoromethyl, nitro, oxo, or combinations thereof, wherein in the alkyl portion one or more $-\text{CH}_2\text{CH}_2-$ groups are each optionally replaced by $-\text{CH}=\text{CH}-$ or $-\text{C}\equiv\text{C}-$, and one or more $-\text{CH}_2-$ groups are each optionally replaced by

-O- or -NH- and/or the alkyl portion is optionally substituted by halogen, oxo, hydroxy, cyano, or combinations thereof;

R^3 is H,

aryl having 6 to 14 carbon atoms and which is unsubstituted or substituted one or more times by halogen, alkyl, alkenyl, alkynyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, OCF_3 , amino, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, 2(-heterocycle)tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxycarbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, trialkylsilyloxy, R^4 -L-, or combinations thereof, or

heteroaryl having 5 to 10 ring atoms in which at least 1 ring atom is a heteroatom, which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxycarbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, trialkylsilyloxy, R^4 -L-, or combinations thereof;

R^4 is H,

alkyl having 1 to 8 carbon atoms, which is unsubstituted or substituted one or more times with halogen, C_{1-4} -alkyl, C_{1-4} -alkoxy, oxo, or combinations thereof,

alkylamino or dialkylamino wherein each alkyl portion has independently 1 to 8 carbon atoms,

a partially unsaturated carbocycle-alkyl group wherein the carbocyclic portion has 5 to 14 carbon atoms and the alkyl portion has 1 to 5 carbon atoms, which is unsubstituted or substituted, one or more times by halogen, alkyl, alkoxy, nitro, cyano, oxo, or combinations thereof,

cycloalkyl having 3 to 10 carbon atoms, which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkoxy, alkyl having 1 to 4 carbon atoms, or combinations thereof,

cycloalkylalkyl having 4 to 16 carbon atoms, which is unsubstituted or substituted in the cycloalkyl portion and/or the alkyl portion one or more times by halogen, oxo, cyano, hydroxy, alkyl, alkoxy or combinations thereof,

aryl having 6 to 14 carbon atoms and which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxycarbonyl, cyano, acyl, alkylthio, alkylsulfinyl, or alkylsulfonyl,

arylalkyl having 7 to 19 carbon atoms, wherein the aryl portion has 6 to 14 carbon atoms and the alkyl portion, which is branched or unbranched, has 1 to 5 carbon atoms, arylalkyl radical is unsubstituted or substituted, in the aryl portion, one or more times by halogen, trifluoromethyl, CF_3O , nitro, amino, alkyl, alkoxy, amino, alkylamino, or dialkylamino and/or substituted in the alkyl portion by halogen, cyano, or methyl,

a heterocyclic group, which is saturated, partially saturated or unsaturated, having

5 to 10 ring atoms in which at least 1 ring atom is a N, O or S atom, which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, or combinations thereof, or

a heterocycle-alkyl group, wherein the heterocyclic portion is saturated, partially saturated or unsaturated, and has 5 to 10 ring atoms in which at least 1 ring atom is a N, O or S atom, and the alkyl portion which is branched or unbranched and has 1 to 5 carbon atoms, the heterocycle-alkyl group is unsubstituted or substituted one or more times in the heterocyclic portion by halogen, alkyl, alkoxy, cyano, trifluoromethyl, CF_3O , nitro, oxo, amino, alkylamino, dialkylamino, or combinations thereof and/or substituted in the alkyl portion by halogen, cyano, or methyl or combinations thereof;

L is a single bond or a divalent aliphatic radical having 1 to 8 carbon atoms wherein one or more $-\text{CH}_2-$ groups are each optionally replaced by $-\text{O}-$, $-\text{S}-$, $-\text{NR}^5-$, $-\text{SO}_2\text{NH}-$, $-\text{NHSO}_2-$, $-\text{CO}-$, $-\text{NR}^5\text{CO}-$, $-\text{CONR}^5-$, $-\text{NHCONH}-$, $-\text{OCONH}-$, $-\text{NHCOO}-$, $-\text{SCONH}-$, $-\text{SCSNH}-$, or $-\text{NHCSNH}-$;

R^5 is H,

alkyl having 1 to 8 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times with halogen, C_{1-4} -alkyl, C_{1-4} -alkoxy, oxo, or combinations thereof;

R^7 is H,

alkyl having 1 to 8 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen,

cycloalkyl having 3 to 10 carbon atoms, which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, or combinations thereof, or

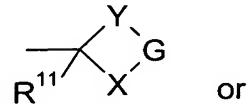
a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is a N, O or S atom, which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxycarbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, or combinations thereof;

R⁸ is H, or

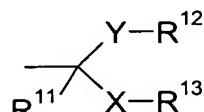
alkyl having 1 to 4 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, cyano, and/or C₁₋₄-alkoxy, and one or more -CH₂CH₂- groups can be replaced in each case by -CH=CH- or -C≡C-;

R⁹ is alkoxy or alkylthio, in each case having 1 to 4 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen;

R^{10} is $-CO-C_{1-4}\text{-alkyl}$ which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, or is



or



;

R^{11} is H or alkyl having 1 to 4 carbon atoms, which is branched or unbranched, and which is unsubstituted or substituted one or more times by halogen;

R^{12} is alkyl having 1 to 6 carbon atoms, which is branched or unbranched, and which is unsubstituted or substituted one or more times by halogen;

R^{13} is alkyl having 1 to 6 carbon atoms, which is branched or unbranched, and which is unsubstituted or substituted one or more times by halogen;

X and Y are each independently O or S; and

G is alkylene having 2 to 7 carbon atoms which is unsubstituted or substituted one or more times by halogen.

66. (Previously Presented): An isolated compound according to claim 1, wherein said compound is selected from:

3'-Chloro-3-cyclopentyloxy-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,

3'-Chloro-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)-3-(3-tetrahydrofuryloxy)diphenylamine,

3'-Cyano-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)-3-((3*R*)-tetrahydrofuryloxy)diphenylamine,

4-Difluoromethoxy-*N*-(1-oxy-3-pyridylmethyl)-3-(3-tetrahydrofuryloxy)diphenylamine,

3,4-Bis(difluoromethoxy)-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,
4-Difluoromethoxy-*N*-(1-oxy-3-pyridylmethyl)-3-((3*R*)-tetrahydrofuryloxy)diphenylamine,
3'-Cyano-4-difluoromethoxy-*N*-(1-oxy-3-pyridylmethyl)-3-((3*R*)-
tetrahydrofuryloxy)diphenylamine,
3'-Chloro-4-difluoromethoxy-*N*-(1-oxy-3-pyridylmethyl)-3-((3*R*)-
tetrahydrofuryloxy)diphenylamine,
4'-*tert*-Butyldimethylsilyloxy-3-cyclopentyloxy-4-methoxy-*N*-(1-oxy-3-
pyridylmethyl)diphenylamine,
N-(3-Cyclopentyloxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic acid,
N-(3-Cyclopentyloxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)-4-aminobenzoic acid,
N-(3-Cyclopentyloxy-4-difluoromethoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic
acid,
N-[4-Methoxy-3-(3-tetrahydrofuryloxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic acid,
N-3,4-Bis(difluoromethoxy)phenyl)-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic acid,
N-[4-methoxy-3-((3*R*)-tetrahydrofuryloxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic
acid,
N-(3-Cyclopropylmethoxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)-4-aminobenzoic acid,
N-(3-Cyclopropylmethoxy-4-difluoromethoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)-3-
aminobenzoic acid,
N-[3-(4-Chlorophenyl)prop-1-yloxy-4-methoxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)-3-
aminobenzoic acid,
N-(3-Cyclopropylmethoxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic acid,
N-[3-(2-Indanyloxy)-4-methoxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic acid,
N-[4-Methoxy-3-(3-tetrahydrofuryloxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic acid,
N-[4-Methoxy-3-((3*R*)-tetrahydrofuryloxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic
acid,
N-[3-(2-Methoxymethoxy)-4-methoxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)-3-aminobenzoic acid,
3-Cyclopropylmethoxy-4-difluoromethoxy-*N*-(1-oxy-3-pyridylmethyl)-4'-(2*H*-tetrazol-5-
yl)diphenylamine,

3-Cyclopentyloxy-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)-4'-(2H-tetrazol-5-yl)diphenylamine,
3-Cyclopentyloxy-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)-3'-(2H-tetrazol-5-yl)diphenylamine,
4-Methoxy-*N*-(1-oxy-3-pyridylmethyl)-3-((3*R*)-tetrahydrofuryloxy)-4'-(2H-tetrazol-5-yl)diphenylamine,
3-Cyclopropylmethoxy-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)-4'-(2H-tetrazol-5-yl)diphenylamine,
4-Difluoromethoxy-*N*-(1-oxy-3-pyridylmethyl)-3-((3*R*)-tetrahydrofuryloxy)-4'-(2H-tetrazol-5-yl)diphenylamine,
3-Cyclopentyloxy-4-difluoromethoxy-*N*-(1-oxy-3-pyridylmethyl)-4'-(2H-tetrazol-5-yl)diphenylamine,
3-Cyclopropylmethoxy-4-difluoromethoxy-*N*-(1-oxy-3-pyridylmethyl)-3'-(2H-tetrazol-5-yl)diphenylamine,
Bis-3,4-difluoromethoxy-*N*-(1-oxy-3-pyridylmethyl)-4'-(2H-tetrazol-5-yl)diphenylamine,
N-(3-Cyclopentyloxy-4-methoxyphenyl)-*N*-(3-pyridyl)-*N*-(1-oxy-3-pyridylmethyl)amine,
N-(3-Cyclopentyloxy-4-difluoromethoxyphenyl)-*N*-(3-pyridyl)-*N*-(1-oxy-3-pyridylmethyl)amine,
N-(3-Cyclopropylmethoxy-4-difluoromethoxyphenyl)-*N*-(3-pyridyl)-*N*-(1-oxy-3-pyridylmethyl)amine,
N-(4-Difluoromethoxy-3-(3*R*)-tetrahydrofuryloxyphenyl)-*N*-(3-pyridyl)-*N*-(1-oxy-3-pyridylmethyl)amine,
3-Cyclopentyloxy-3'-ethanesulfonylamino-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,
3-Cyclopentyloxy-4-methoxy-3'-(1-propanesulfonylamino)-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,
3-Cyclopentyloxy-4'-ethanesulfonylamino-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,
3-Cyclopentyloxy-4-methoxy-4'-(1-propanesulfonylamino)-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,
3-Cyclopropylmethoxy-3'-ethanesulfonylamino-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,
4-Difluoromethoxy-3'-ethanesulfonylamino-*N*-(1-oxy-3-pyridylmethyl)-3-[(3*R*)-tetrahydrofuryloxy]diphenylamine,

4-Methoxy-3-[2-(2-pyridyl)ethoxy]-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,
4-Methoxy-*N*-(1-oxy-3-pyridylmethyl)-3-[(3*R*)-tetrahydrofuryloxy]diphenylamine,
3'-Chloro-4-methoxy-3-[2-(2-pyridyl)ethoxy]-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,
3'-Chloro-4-methoxy-*N*-(1-oxy-3-pyridylmethyl)-3-[(3*R*)-tetrahydrofuryloxy]diphenylamine,
3-Cyclopentyloxy-4-methoxy-4'-[2-(5-oxopyrrolidinyl)methoxy]-*N*-(1-oxy-3-pyridylmethyl)diphenylamine,
3-Cyclopentyloxy-4-methoxy-*N*-(3-aminocarbonylphenyl)-*N*-(1-oxy-3-pyridylmethyl)aniline,
3,4-Bisdifluoromethoxy-*N*-(3-carboxy-4-chlorophenyl)-*N*-(1-oxy-3-pyridylmethyl)aniline,
3,4-Bisdifluoromethoxy-*N*-(4-(1-pyrrol-1-yl)phenyl)-*N*-(1-oxy-3-pyridylmethyl)aniline,
4-Methoxy-3-(*R*)-tetrahydrofuryloxy-*N*-(3-carboxy-4-chlorophenyl)-*N*-(1-oxy-3-pyridylmethyl)aniline,
4-Methoxy-3-(*R*)-tetrahydrofuryloxy-*N*-(3-carboxyphenyl)-*N*-(1-oxy-4-pyridylmethyl)aniline,
4-Methoxy-3-(*R*)-tetrahydrofuryloxy-*N*-(3-pyridyl)-*N*-(1-oxy-4-pyridylmethyl)aniline,
3-Cyclopentyloxy-4-methoxy-*N*-(4-carboxy-3-chlorophenyl)-*N*-(1-oxy-3-pyridylmethyl)aniline,
2-Acetyl-7-methoxy-4-(*N*-(4-cyanophenyl)-*N*-(1-oxy-3-pyridylmethyl))aminobenzofuran,
2-Acetyl-7-methoxy-4-(*N*-phenyl-*N*-(1-oxy-4-pyridylmethyl))aminobenzofuran,
2-Acetyl-7-methoxy-4-(*N*-(3-carboxyphenyl)-*N*-(1-oxy-3-pyridylmethyl))aminobenzofuran,
1-Cyclopentyl-3-ethyl-6-(*N*-(3-carboxyphenyl)-*N*-(1-oxy-3-pyridylmethyl))aminoindazole,
2-Acetyl-7-methoxy-4-(*N*-(4-acetylphenyl)-*N*-(1-oxy-3-pyridylmethyl))aminobenzofuran,
N-[4-Methoxy-3-((3*R*)-3-tetrahydrofuranyl)oxyphenyl]-4-methylsulfonylaminocarbonyl-*N*-(1-oxy-3-pyridylmethyl)aniline,
4-(4-Fluorophenyl)sulfonylaminocarbonyl-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuranyl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)aniline,
3-Chloro-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuranyl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)aniline,
3-Amino-*N*-(5-fluoro-1-oxy-3-pyridylmethyl)-*N*-(4-methoxy-3-((3*R*)-3-tetrahydrofuranyl)oxyphenyl]benzoic acid,
3-Amino-*N*-(3-cyclopentyloxy-4-methoxyphenyl)-*N*-(1-oxy-2-pyridylmethyl)benzoic acid,
3-Amino-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuranyl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)-5-

trifluoromethylbenzoic acid,

4-Ethylsulfonylaminocarbonyl-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuran-3-yl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)aniline,

4-(2-Fluorophenyl)sulfonylaminocarbonyl-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuran-3-yl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)aniline

4-(3-Chlorophenyl)sulfonylaminocarbonyl-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuran-3-yl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)aniline,

3-Amino-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuran-3-yl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)-6-trifluoromethylbenzoic acid,

4-Amino-*N*-[4-difluoromethoxy-3-((3*R*)-3-tetrahydrofuran-3-yl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,

N-[4-Difluoromethoxy-3-((3*R*)-3-tetrahydrofuran-3-yl)oxyphenyl]-4-methylsulfonylaminocarbonyl-*N*-(1-oxy-3-pyridylmethyl)aniline,

N-[4-Methoxy-3-((3*R*)-3-tetrahydrofuran-3-yl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)-4-phenylsulfonylaminocarbonylaniline,

3-Amino-*N*-(3-cyclopentyloxy-4-methoxyphenyl)-*N*-(5-fluoro-1-oxy-3-pyridylmethyl)benzoic acid,

4-Amino-*N*-(3-cyclopentyloxy-4-methoxyphenyl)-*N*-(5-fluoro-1-oxy-3-pyridylmethyl)benzoic acid,

3-Amino-*N*-[4-difluoromethoxy-3-((3*R*)-3-tetrahydrofuran-3-yl)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,

3-Amino-*N*-(3-cyclobutyloxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,

3-Amino-*N*-(3-cyclopentyloxy-4-methoxyphenyl)-5-fluoro-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,

3-Amino-*N*-[3,4-bis(difluoromethoxy)phenyl]-5-fluoro-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,

4-Amino-*N*-(3-cyclobutyloxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,

4-Amino-*N*-(3-ethoxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,

4-Amino-*N*-(3-isopropoxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,

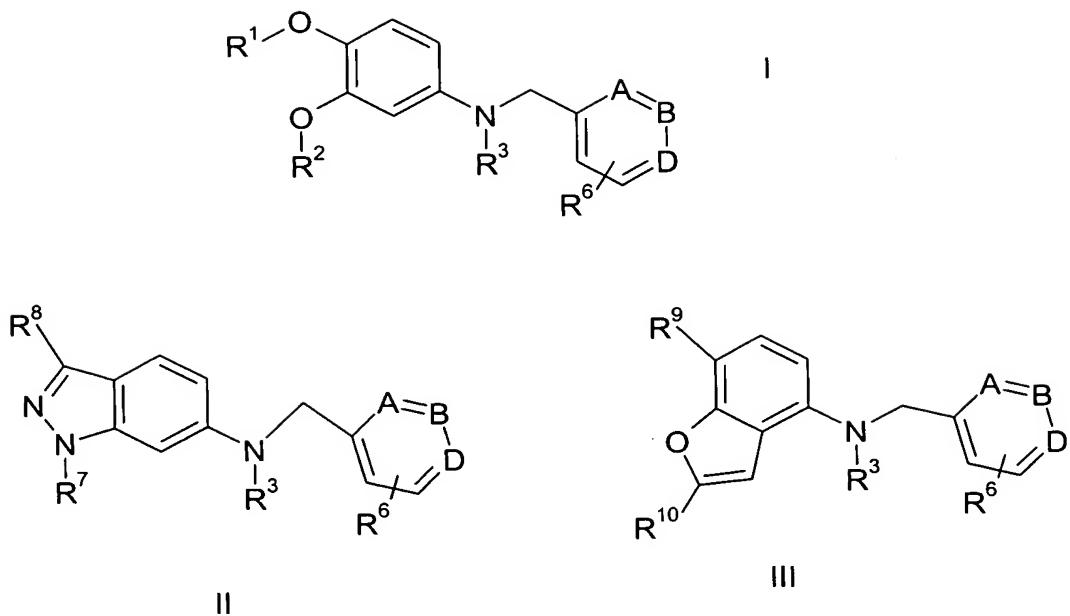
N-[4-Difluoromethoxy-3-((3*R*)-3-tetrahydrofuran-3-yl)oxyphenyl]-4-(3,4-

difluorophenyl)sulfonylaminocarbonyl-*N*-(1-oxy-3-pyridylmethyl)aniline,
4-Amino-*N*-(3-cyclopropylmethoxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
N-[3,4-Bis(difluoromethoxy)phenyl]-4-(4-fluorophenyl)sulfonylaminocarbonyl-*N*-(1-oxy-3-pyridylmethyl)aniline,
4-(2,4-Difluorophenyl)sulfonylaminocarbonyl-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuran-*Y*)-oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)aniline,
4-(3,4-Difluorophenyl)sulfonylaminocarbonyl-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuran-*Y*)-oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)aniline,
N-[4-Difluoromethoxy-3-((3*R*)-3-tetrahydrofuran-*Y*)-oxyphenyl]-4-ethylsulfonylaminocarbonyl-*N*-(1-oxy-3-pyridylmethyl)aniline,
3-Amino-*N*-(3,4-dimethoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
3-Amino-*N*-(3-ethoxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
3-Amino-*N*-(3-isopropoxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
4-(3,4-Difluorophenyl)sulfonylaminocarbonyl-*N*-(3-ethoxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)aniline,
3-Amino-*N*-[3,4-bis(difluoromethoxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
3-Amino-*N*-(4-difluoromethoxy-3-ethoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
4-Amino-*N*-(4-difluoromethoxy-3-ethoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
3-Amino-*N*-(4-difluoromethoxy-3-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
4-Amino-*N*-(3-cyclopentyloxy-4-methoxyphenyl)-*N*-(1-oxypyridin-3-ylmethyl)pyridine,
N-[Bis-3,4-(difluoromethoxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)-4-[2-(2-tetrahydropyran-*Y*)-2H-tetrazol-5-yl]aniline,
N-[Bis-3,4-(difluoromethoxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)-3-(2H-tetrazol-5-yl)aniline; and
pharmaceutically acceptable salts thereof,
wherein optically active compounds can be in the form of their separate enantiomers or mixtures
thereof, including racemic mixtures.

67. (Previously Presented): An isolated compound according to claim 1, wherein said compound is selected from:

4-amino-*N*-(3-cyclopentyloxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)pyridine, *N*-[Bis-3,4-(difluoromethoxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)-4-[2-(2-tetrahydropyranyl)-2H-tetrazol-5-yl]aniline,
3-Chloro-*N*-[4-methoxy-3-((3*R*)-3-tetrahydrofuran-*R*)oxyphenyl]-*N*-(1-oxy-3-pyridylmethyl)aniline,
4-Amino-*N*-(3-cyclopentyloxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
3-Amino-*N*-(3-cyclopentyloxy-4-methoxyphenyl)-*N*-(1-oxy-3-pyridylmethyl)benzoic acid,
3-Amino-*N*-[bis-3,4-(difluoromethoxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)benzoic acid, *N*-[Bis-3,4-(difluoromethoxy)phenyl]-*N*-(1-oxy-3-pyridylmethyl)-3-(2H-tetrazol-5-yl)aniline; and
pharmaceutically acceptable salts thereof,
wherein optically active compounds can be in the form of their separate enantiomers or mixtures thereof, including racemic mixtures.

68. (Previously Presented): A pharmaceutical composition containing a pharmaceutically acceptable carrier and a compound of Formulas I-III:



wherein

one of A, B and D is N-O and the others are CR⁶;

R¹ is alkyl having 1 to 4 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen;

R² is alkyl having 1 to 12 carbon atoms which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, hydroxy, cyano, C₁-₄-alkoxy, oxo or combinations thereof, and wherein optionally one or more -CH₂CH₂- groups is replaced in each case by -CH=CH- or -C≡C-,

cycloalkyl having 3 to 10 carbon atoms which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, or combinations thereof,

cycloalkylalkyl having 4 to 16 carbon atoms which is unsubstituted or substituted

in the cycloalkyl portion and/or the alkyl portion one or more times by halogen, oxo, cyano, hydroxy, C₁₋₄-alkyl, C₁₋₄-alkoxy or combinations thereof,

aryl having 6 to 14 carbon atoms, which is unsubstituted or substituted one or more times by halogen, CF₃, OCF₃, alkyl, hydroxy, alkoxy, nitro, methylenedioxy, ethylenedioxy, cyano, or combinations thereof,

arylalkyl in which the aryl portion has 6 to 14 carbon atoms and the alkyl portion, which is branched or unbranched, has 1 to 5 carbon atoms, wherein the arylalkyl radical is unsubstituted or is substituted in the aryl portion one or more times by halogen, CF₃, OCF₃, alkyl, hydroxy, alkoxy, nitro, cyano, methylenedioxy, ethylenedioxy, or combinations thereof, and wherein in the alkyl portion one or more -CH₂CH₂- groups are each optionally replaced by -CH=CH- or -C≡C-, and/or one or more -CH₂- groups are each optionally replaced by -O- or -NH- and/or the alkyl portion is optionally substituted by halogen, oxo, hydroxy, cyano, or combinations thereof,

a partially unsaturated carbocyclic group having 5 to 14 carbon atoms, which is unsubstituted or substituted one or more times by halogen, alkyl, alkoxy, hydroxy, nitro, cyano, oxo, or combinations thereof,

a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is an N, O or S atom, wherein the heterocyclic group is unsubstituted or substituted one or more times by halogen, hydroxy, aryl, alkyl, alkoxy, cyano, trifluoromethyl, nitro, oxo, or combinations thereof, or

a heterocycle-alkyl group, wherein the heterocyclic portion is saturated, partially saturated or unsaturated, and has 5 to 10 ring atoms in which at least 1 ring atom

is an N, O or S atom, and the alkyl portion is branched or unbranched and has 1 to 5 carbon atoms, the heterocycle-alkyl group is unsubstituted or substituted one or more times in the heterocyclic portion by halogen, OCF₃, hydroxy, aryl, alkyl, alkoxy, cyano, trifluoromethyl, nitro, oxo, or combinations thereof, wherein in the alkyl portion one or more -CH₂CH₂- groups are each optionally replaced by -CH=CH- or -C≡C-, and/or one or more -CH₂- groups are each optionally replaced by -O- or -NH- and/or the alkyl portion is optionally substituted by halogen, oxo, hydroxy, cyano, or combinations thereof;

R³ is H,

cycloalkyl having 3 to 10 carbon atoms which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, or combinations thereof,

aryl having 6 to 14 carbon atoms and which is unsubstituted or substituted one or more times by halogen, alkyl, alkenyl, alkynyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, OCF₃, amino, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, pyrrolyl, tetrazole-5-yl, 2(-heterocycle)tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxycarbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, trialkylsilyloxy, R⁴-L-, or combinations thereof,

heteroaryl having 5 to 10 ring atoms in which at least 1 ring atom is a heteroatom, which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxycarbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy,

trialkylsilyloxy, R^4 -L-, or combinations thereof, or

a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is an N, O or S atom, which is unsubstituted or substituted one or more times by halogen, hydroxy, aryl, alkyl, alkoxy, cyano, trifluoromethyl, nitro, oxo, or combinations thereof;

R^4 is H,

alkyl having 1 to 8 carbon atoms which is unsubstituted or substituted one or more times by halogen, C_{1-4} -alkyl, C_{1-4} -alkoxy, oxo, or combinations thereof,

alkylamino or dialkylamino wherein each alkyl portion has independently 1 to 8 carbon atoms,

a partially unsaturated carbocycle-alkyl group wherein the carbocyclic portion has 5 to 14 carbon atoms and the alkyl portion has 1 to 5 carbon atoms, which is unsubstituted or substituted one or more times by halogen, alkyl, alkoxy, nitro, cyano, oxo, or combinations thereof,

cycloalkyl having 3 to 10 carbon atoms which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkoxy, alkyl having 1 to 4 carbon atoms, or combinations thereof,

cycloalkylalkyl having 4 to 16 carbon atoms which is unsubstituted or substituted in the cycloalkyl portion and/or the alkyl portion one or more times by halogen, oxo, cyano, hydroxy, alkyl, alkoxy or combinations thereof,

aryl having 6 to 14 carbon atoms and which is unsubstituted or substituted one or

more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy, dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, cycloalkyl, aryl, heteroaryl, or combinations thereof,

arylalkyl having 7 to 19 carbon atoms, wherein the aryl portion has 6 to 14 carbon atoms and the alkyl portion, which is branched or unbranched, has 1 to 5 carbon atoms, wherein the arylalkyl radical is unsubstituted or substituted, in the aryl portion, one or more times by halogen, trifluoromethyl, CF_3O , nitro, amino, alkyl, alkoxy, amino, alkylamino, dialkylamino, or combinations thereof, and/or substituted in the alkyl portion by halogen, cyano, methyl, or combinations thereof, wherein in the alkyl portion one or more $-\text{CH}_2\text{CH}_2-$ groups are each optionally replaced by $-\text{CH}=\text{CH}-$ or $-\text{C}\equiv\text{C}-$, and/or one or more $-\text{CH}_2-$ groups are each optionally replaced by $-\text{O}-$ or $-\text{NH}-$,

a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is an N, O or S atom, which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, or combinations thereof, or

a heterocycle-alkyl group, wherein the heterocyclic portion is saturated, partially saturated or unsaturated, and has 5 to 10 ring atoms in which at least 1 ring atom is a N, O or S atom, and the alkyl portion which is branched or unbranched and has 1 to 5 carbon atoms, the heterocycle-alkyl group is unsubstituted or

substituted one or more times in the heterocyclic portion by halogen, alkyl, alkoxy, cyano, trifluoromethyl, CF_3O , nitro, oxo, amino, alkylamino, dialkylamino, or combinations thereof and/or substituted in the alkyl portion by halogen, cyano, or methyl or combinations thereof;

L is a single bond or a divalent aliphatic radical having 1 to 8 carbon atoms wherein one or more $-\text{CH}_2-$ groups are each optionally replaced by $-\text{O}-$, $-\text{S}-$, $-\text{SO}-$, $-\text{SO}_2-$, $-\text{NR}^5-$, $-\text{SO}_2\text{NH}-$, $-\text{NHSO}_2-$, $-\text{SO}_2\text{NR}^5-$, $-\text{NR}^5\text{SO}_2-$, $-\text{CO}-$, $-\text{NR}^5\text{CO}-$, $-\text{CONR}^5-$, $-\text{NHCONH}-$, $-\text{OCONH}-$, $-\text{NHCOO}-$, $-\text{SCONH}-$, $-\text{SCSNH}-$, or $-\text{NHCSNH}-$;

R^5 is H,

alkyl having 1 to 8 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, C_{1-4} -alkyl, C_{1-4} -alkoxy, oxo, or combinations thereof,

aryl having 6 to 14 carbon atoms and which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, ethylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, or combinations thereof, or

arylalkyl having 7 to 19 carbon atoms, wherein the aryl portion has 6 to 14 carbon atoms and the alkyl portion, which is branched or unbranched, has 1 to 5 carbon atoms, wherein the arylalkyl radical is unsubstituted or substituted, in the aryl portion, one or more times by halogen, trifluoromethyl, CF_3O , nitro, amino, alkyl,

alkoxy, amino, alkylamino, dialkylamino, or combinations thereof, and/or substituted in the alkyl portion by halogen, cyano, methyl, or combinations thereof, wherein in the alkyl portion one or more -CH₂CH₂- groups are each optionally replaced by -CH=CH- or -C≡C-, and/or one or more -CH₂- groups are each optionally replaced by -O- or -NH-;

R⁶ is H, halogen, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, CN, or hydroxyl;

R⁷ is H,

alkyl having 1 to 8 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, cyano, hydroxy, C₁₋₄-alkoxy, or combinations thereof

cycloalkyl having 3 to 10 carbon atoms which is unsubstituted or substituted one or more times by halogen, hydroxy, oxo, cyano, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, or combinations thereof,

cycloalkylalkyl having 4 to 16 carbon atoms which is unsubstituted or substituted in the cycloalkyl portion and/or the alkyl portion one or more times by halogen, oxo, cyano, hydroxy, C₁₋₄-alkyl, C₁₋₄-alkoxy or combinations thereof,

aryl having 6 to 14 carbon atoms, which is unsubstituted or substituted one or more times by halogen, CF₃, OCF₃, alkyl, hydroxy, alkoxy, nitro, methylenedioxy, ethylenedioxy, cyano, or combinations thereof,

arylalkyl in which the aryl portion has 6 to 14 carbon atoms and the alkyl portion,

which is branched or unbranched, has 1 to 5 carbon atoms, wherein the arylalkyl radical is unsubstituted or is substituted in the aryl portion one or more times by halogen, CF_3 , OCF_3 , alkyl, hydroxy, alkoxy, nitro, cyano, methylenedioxy, ethylenedioxy, or combinations thereof, and wherein in the alkyl portion one or more $-\text{CH}_2\text{CH}_2-$ groups are each optionally replaced by $-\text{CH}=\text{CH}-$ or $-\text{C}\equiv\text{C}-$, and/or one or more $-\text{CH}_2-$ groups are each optionally replaced by $-\text{O}-$ or $-\text{NH}-$ and/or the alkyl portion is optionally substituted by halogen, oxo, hydroxy, cyano, or combinations thereof,

a heterocyclic group, which is saturated, partially saturated or unsaturated, having 5 to 10 ring atoms in which at least 1 ring atom is an N, O or S atom, which is unsubstituted or substituted one or more times by halogen, alkyl, hydroxy, alkoxy, alkoxyalkoxy, nitro, methylenedioxy, ethylenedioxy, trifluoromethyl, amino, aminomethyl, aminoalkyl, aminoalkoxy dialkylamino, hydroxyalkyl, hydroxamic acid, tetrazole-5-yl, hydroxyalkoxy, carboxy, alkoxy carbonyl, cyano, acyl, alkylthio, alkylsulfinyl, alkylsulfonyl, phenoxy, or combinations thereof, or

a heterocycle-alkyl group, wherein the heterocyclic portion is saturated, partially saturated or unsaturated, and has 5 to 10 ring atoms in which at least 1 ring atom is a N, O or S atom, and the alkyl portion which is branched or unbranched and has 1 to 5 carbon atoms, the heterocycle-alkyl group is unsubstituted or substituted one or more times in the heterocyclic portion by halogen, alkyl, alkoxy, cyano, trifluoromethyl, CF_3O , nitro, oxo, amino, alkylamino, dialkylamino, or combinations thereof and/or substituted in the alkyl portion by halogen, cyano, or methyl or combinations thereof;

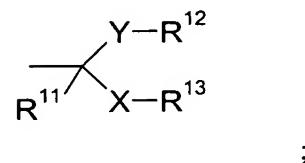
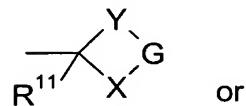
R^8 is H, or

alkyl having 1 to 4 carbon atoms, which is branched or unbranched and which is

unsubstituted or substituted one or more times by halogen, cyano, and/or C₁₋₄-alkoxy, and one or more -CH₂CH₂- groups can be replaced in each case by -CH=CH- or -C≡C-;

R⁹ is alkoxy or alkylthio, in each case having 1 to 4 carbon atoms, which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen;

R¹⁰ is -CO-C₁₋₄-alkyl which is branched or unbranched and which is unsubstituted or substituted one or more times by halogen, or is



R¹¹ is H or alkyl having 1 to 4 carbon atoms, which is branched or unbranched, and which is unsubstituted or substituted one or more times by halogen;

R¹² is alkyl having 1 to 6 carbon atoms, which is branched or unbranched, and which is unsubstituted or substituted one or more times by halogen;

R¹³ is alkyl having 1 to 6 carbon atoms, which is branched or unbranched, and which is unsubstituted or substituted one or more times by halogen;

X and Y are each independently O or S; and

G is alkylene having 2 to 7 carbon atoms which is unsubstituted or substituted one or more times by halogen; or
a pharmaceutically acceptable salt thereof;

wherein an optically active compound can be in the form of one of its separate enantiomers or mixtures thereof, including racemic mixtures.

69. (Previously Presented): A composition of claim 68, wherein the compound is provided in a unit dosage of 0.1 - 50 mg.

70. (Currently Amended): A method for enhancing cognition and/or treating psychosis in a patient comprising administering to said patient an effective amount of a composition according to claim 68.

wherein

~~one of A, B and D is N or O and the others are CR⁶;~~

71. (Original): A method according to claim 70, wherein said compound is administered in an amount of 0.01-100 mg/kg of body weight/day.

72. (Original): A method according to claim 70, wherein said patient is a human.

73. (Original): A method of claim 70, wherein the patient is suffering from cognition impairment or decline.

74. (Original): A method according to claim 70, wherein said patient is suffering

from memory impairment.

75. (Currently Amended): A method according to claim 74, wherein said patient is suffering from memory impairment due to Alzheimer's disease, multiple sclerosis, amyolaterosclerosis, multiple systems atrophy, schizophrenia, Parkinson's disease, Huntington's disease, Pick's disease, Creutzfeld-Jakob disease, depression, aging, head trauma, stroke, spinal cord injury, CNS hypoxia, cerebral senility, diabetes associated cognitive impairment, memory deficits from early exposure of anesthetic agents, multiinfarct dementia, an acute neuronal disease, HIV, cardiovascular disease, or age-related cognitive decline.

76. (Original): A method according to claim 74, wherein said patient is suffering from memory impairment due to dementia.

77. (Original): A method according to claim 70, wherein said patient is suffering from a psychosis.

78. (Currently Amended): The method of claim 77, wherein the psychosis is schizophrenia, bipolar or manic depression, or major depression, ~~drug addiction or morphine dependence~~.

79. (Cancelled):

80. (Original): A method according to claim 70, wherein the patient is treated to effect PDE4 enzyme inhibition.

81. (Previously Presented): A method of treating a patient suffering from an allergic or inflammatory disease comprising administering to said patient an effective amount of a composition according to claim 68.

82. (Previously Presented): A method of treating a patient suffering from neurodegeneration resulting from a disease or injury comprising administering to said patient an effective amount of a composition according to claim 68.

83. (Original): The method of claim 82, wherein the disease or injury is stroke, spinal cord injury, Alzheimer's disease, multiple sclerosis, amyotrophic lateral sclerosis (ALS), or multiple systems atrophy (MSA).

84. (Cancelled):

85. (Previously Presented): An isolated compound according to claim 1, wherein said compound is in the form of a substantially pure enantiomer.

86. (New): A method for treating drug addiction or morphine dependence in a patient comprising administering to said patient an effective amount of a composition according to claim 68.